



09/674733

C of C

IN OFFICE UNITED STATES PATENT AND TRADEMARK OFFICE

Atty. Docket: SZARDENINGS=1

| | | |
|------------------------------|---|----------------------|
| In re Patent of: |) | Conf. No.: 3759 |
| |) | |
| SZARDENINGS et al. |) | |
| |) | |
| Patent No.: 7,008,925 |) | Washington, D.C. |
| |) | |
| Issued: March 7, 2006 |) | April 20, 2007 |
| |) | |
| For: MELANOCORTIN 1 RECEPTOR |) | |
| SELECTIVE COMPOUNDS |) | ATTN: Certificate of |
| |) | Correction Division |

REQUEST FOR EXPEDITED CERTIFICATE OF CORRECTION UNDER 37 C.F.R.
§1.322

Honorable Commissioner for Patents
U.S. Patent and Trademark Office
Randolph Building, Mail Stop Post Issue
401 Dulany Street
Alexandria, VA 22314

Certificate

APR 24 2007

of Correction

Sir:

In checking over the printed copy of the above-identified patent, we have found the following error that is the fault of the Patent and Trademark Office. It is respectfully requested that this error be corrected in accordance with 37 CFR §1.322(a). The error to be corrected is listed below.

The PTO erred by publishing, as the sequence listing for this case (cols. 39-46), a sequence listing associated with some other case. This is evident from comparison of cols. 39-46 with the 8 page sequence listing filed June 12, 2001.

APR 25 2007

In re of U.S. Patent 7,008,925

| | <u>Cols. 39-46</u> | <u>June 12, 2001</u> |
|---------------------|-------------------------|---------------------------------|
| Number of Sequences | 22 | 15 |
| SEQ ID NO: 1 | 30 base DNA sequence | 13 a.a. Polypeptide sequence |
| SEQ ID NO: 2 | 10 base DNA sequence | 13 a.a. Polypeptide sequence |

and so forth.

It is important that the error be corrected because one or more of SEQ ID NOS: 1-15 are referred to in claims 9 or 10.

We are attaching one copy of the Certificate of Correction form.


In accordance with MPEP §1480.01, in an effort to expedite processing of this request, copies of cols. 39-46 of the patent, and the June 12, 2001, filing are attached, so that the PTO can verify that an error was made. Consequently, correction should be expedited pursuant to MPEP §1480.01.

Granting of this request is earnestly solicited.

Respectfully submitted,

BROWDY AND NEIMARK, P.L.L.C.
Attorneys for Applicant(s)

By


Iver P. Cooper
Registration No. 28,005

:dtb

Telephone No.: (202) 628-5197

Facsimile No.: (202) 737-3528

G:\ipc\n-q\Plou\Szardenings1\2007-04-20CertCor322PTOFault.doc

APR 25 2007

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

Page 1 of 13

PATENT NO. : 7,008,925
APPLICATION NO.: 09/674,733
ISSUE DATE : March 7, 2006
INVENTOR(S) : SZARDENINGS et al.

It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Please delete the sequence listing at cols. 39-46 and insert therefor the following sequence listing:

SEQUENCE LISTING

<110> Szardenings, Michael

Muceniece, Ruta

Mutule, Ilze

Mutulis, Felikss

Jarl, Wikberg

<120> Melanocortin 1 Receptor Selective Compounds

<130> 1085.0050000/RWE/ALS

<140> 09/674,733

<141> 1999-05-05

MAILING ADDRESS OF SENDER (Please do not use customer number below):

624 Ninth Street, NW
Suite 300
Washington, DC 20001-5303

This collection of information is required by 37 CFR 1.322, 1.323, and 1.324. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 1.0 hour to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: **Attention Certificate of Corrections Branch, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

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**UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION**Page 2 of 13

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APPLICATION NO.: 09/674,733
ISSUE DATE : March 7, 2006
INVENTOR(S) : SZARDENINGS et al.

It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

<150> PCT/GB99/01388

<151> 1999-05-05

<150> SE 9801571-2

<151> 1998-05-05

<160> 15

<170> PatentIn version 3.0

<210> 1

<211> 13

<212> PRT

<213> Artificial

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It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

<220>

<223> Synthetic peptide with high affinity for melanocortin receptor 1

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Ser Ser Ile Ile Ser His Phe Arg Trp Gly Lys Pro Val
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<223> D amino acid

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<211> 14

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Tyr Ser Ser Ile Ile Ser His Phe Arg Trp Gly Lys Pro Val
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<210> 4

<211> 13

<212> PRT

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<213> Artificial
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<223> Synthetic peptide with high affinity for melanocortin receptor 1

<400> 4

Tyr Ser Ile Ile Ser His Phe Arg Trp Gly Lys Pro Val
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<212> PRT

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Ser Ser Ile Ile Ser His Phe Arg Trp Gly Lys Pro Val Tyr
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<211> 13

<212> PRT

<213> Artificial

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It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

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<210> 9

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<211> 13

<212> PRT

<213> Artificial

<220>

<223> Synthetic peptide with high affinity for melanocortin receptor 1

<400> 9

| | | | | | | | | | | | | |
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It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

<211> 13

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<213> Artificial

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<223> Synthetic peptide with high affinity for melanocortin receptor 1

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<222> (7)..(7)

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Ser Ser Ile Ile Ser His Phe Arg Trp Gly Lys Pro Val
1 5 10

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2 5 2007

Cell Culture

RAW 264.7 cells (TIB-71), obtained from American Type Culture Collection, 12301 Parklawn Drive, Rockville, Md., 20852, USA, and cultured in Dulbecco's modified Eagle medium (Gibco, BRL) supplemented with 10% heat-inactivated fetal bovine serum, 100 IU penicillin/ml and 100 µg streptomycin/ml at 37° C. in a humidified atmosphere of 95% air and 5% CO₂. Cells grown in monolayers were detached from the culture flasks and collected by low speed centrifugation (700×g).

Incubation of Compounds of the Invention with RAW 264.7 Cells

The cells obtained as above were resuspended in F-12 (HAM) medium (Gibco, BRL) and distributed into 96-well plates at a density of 2.5×10⁶ cells per well, and incubated with 100 ng/mL bacterial lipopolysaccharide (L4391, Sigma Chemical Company, P.O. Box 14508, St. Louis, Mo. 63178, USA), 5 units/mL of mouse recombinant interferon gamma (15517, Sigma Chemical Company, P.O. Box 14508, St. Louis, Mo. 63178, USA) and the compounds of the invention using concentrations ranging 01 µM, for 16 h, whereafter an aliquot of the medium was collected for measurement of nitric oxide (NO).

Measurement of Nitric Oxide

Nitric oxide was measured by monitoring the nitrite production essentially using the method of Wishnok et al. (Methods in Enzymology, 1996, 268, 130–151). In brief 50 µL of culture medium was mixed with 50 µL Griess reagent (i.e. a 1:1 mixture of 0.1% N-naphthylethylenediamine dihydrochloride and 1% sulfanilamide in 5% (v/v) phosphoric acid) and after 10 min the absorption was measured at 540 nm. The nitrite concentrations were calculated from a standard curve constructed, by instead of culture medium, adding 50 µL of between 3 to 100 µM of NaNO₂ to the assays.

Results

The results are shown in FIG. 6. As can be seen from the Figure, the compounds of the invention, MSO5 and MSO9, as well as α-MSH caused a strong dose dependent inhibition of the NO-production, the potencies and efficacies of MSO5 and MSO9 being similar to that of the α-MSH. This data shows that MSO5 and MSO9 share the capacity of α-MSH to inhibit inflammation. This is because NO is a key component of inflammation.

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COPY

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE



In re application of:

Szardenings *et al.*

Appl. No. 09/674,733
(U.S. Natl. Phase of PCT/GB99/01388)

Int'l Filing Date: May 5, 1999

For: **Melanocortin 1 Receptor
Selective Compounds**

Art Unit: *To be assigned*

Examiner: *To be assigned*

Atty. Docket: 1085.0050000/RWE/ALS

Amendment and Submission of Substitute Sequence Listing Under 37 C.F.R. § 1.825(a)

Commissioner for Patents
Washington, D.C. 20231

Sir:

In compliance with 37 C.F.R. § 1.825(a), Applicants submit substitute sheets to amend the paper copy of the Sequence Listing.

In the Specification:

Please cancel the existing Sequence Listing for the above-identified application, replace it with the substitute Sequence Listing appended hereto, and insert the same at the end of the application.

Remarks

Applicants' Agent hereby states that the change made in the sequence listing does not include new matter. Applicants' undersigned Agent has amended the specification only to direct the entry of this corrected Sequence Listing at the end of the application.

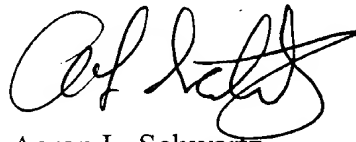
APR 25 2007

In accordance with 37 C.F.R. § 1.825(b), the paper copy of the Sequence Listing and the computer readable copy of the Sequence Listing submitted herewith are the same.

It is respectfully believed this application is now in condition for allowance. Early notice to this effect is earnestly solicited.

Respectfully submitted,

STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.



Aaron L. Schwartz
Agent for Applicants
Provisional Registration No. P-48,181

Date: 6/12/01

1100 New York Avenue, N.W.
Suite 600
Washington, D.C. 20005-3934
(202) 371-2600

P:\USERS\Schwartz\Cases\1085\005\pto\825 amendment.wpd

APR 25 2007

SEQUENCE LISTING



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SEP 25 2007